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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : C09C 1/30, C09D 7/00	A1	(11) International Publication Number: WO 96/34062 (43) International Publication Date: 31 October 1996 (31.10.96)
(21) International Application Number: PCT/EP96/01682 (22) International Filing Date: 23 April 1996 (23.04.96) (30) Priority Data: 195 16 253.6 26 April 1995 (26.04.95) DE (71) Applicant (for all designated States except US): GRACE GMBH [DE/DE]; Erlengang 31, D-22844 Norderstedt (DE). (72) Inventors; and (75) Inventors/Applicants (for US only): BELLIGOI, Peter [DE/DE]; In den Weingärten 28, D-67551 Worms (DE). FIELD, Rex [DE/DE]; Westhofener Strasse 29, D-67596 Dittelsheim-Hessloch (DE). LÜERS, Georg [DE/DE]; Bergstrasse 23, D-67593 Westhofen (DE). SCHNEIDER, Rita [DE/DE]; Kurt-Schumacher-Strasse 18, D-67551 Worms (DE). (74) Agent: VAN HEESCH, Helmut; Uexküll & Stolberg, Beselerstrasse 4, D-22607 Hamburg (DE).		(81) Designated States: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: MATTING AGENT BASED ON AGGREGATED SILICA		
(57) Abstract The invention relates to an aggregated silica gel which has been produced from silica gel particles having a particle size of 1 to 20 μ , a surface of 200 to 1000 m ² /g and a specific pore volume of 0.4 to 2.5 ml/g and a binding agent selected from synthetic or natural phyllosilicate, pyrogenic silicon dioxide and organic polymers soluble in water or dispersible in water. Aggregation takes place by spray-drying a suspension having 1 to 25 % solids. The product is suitable as a matting agent for coatings, as an antiblocking agent for polymer films and as a beer clarifying agent.		

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Matting agent based on aggregated silica

Modern matting agents for coatings consist of highly porous, amorphous silicon dioxide which has been adjusted to a particle size which is appropriate for the required coating film thickness.

The most widely used matting agents originate from two processes:

- gelation of silicic acid, washing, drying, milling
- precipitation of silicic acid in such a way that a "reinforced" structure is formed, washing, drying, milling.

The best representatives of both processes show a high matting efficiency coupled with low thickening, good dispersibility and lacquer film transparency (clarity). SYLOID ED5, which is manufactured by Grace GmbH, Worms, Germany, is a typical representative of the first class and is used throughout this description as a standard for the state of the art.

Another type of matting agent has also been developed which exhibits an increased matting efficiency compared with the two aforementioned classes. This is attained by agglomerating in various ways a pyrolytic (pyrogenic, fumed) silicon dioxide (silica) (pyrolytic silicon dioxide is obtained by flame hydrolysis of SiCl_4). This type of product (e.g. TS 100 from

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Degussa) exhibits a matting efficiency which is increased by 10 to 25 % compared with the first two types. However, it is a major disadvantage of this type of product that it increases the viscosity of the paint (lacquer) in which it is used, even at the lower concentrations which are required in order to obtain the same gloss as with the other products. This increase in viscosity makes processing of the paint more difficult. A further disadvantage is the reduced dispersibility.

DE 12 09 108 describes the spray-drying of an aqueous suspension containing 20 to 60 % pyrolytic silicon dioxide in order to obtain a spherical product having an average particle size of approx. 200 μm .

DE 24 14 478 describes the production of a matting agent by aggregation of powdery pyrolytic silicon dioxide. The silicon dioxide is wetted by adding approx. 5 % of water, based on silicon dioxide, and then dried in an oven or an air-jet mill. The surface of the powder is coated only with a very thin (monomolecular) layer because of low moisture absorption. During subsequent evaporation, aggregation takes place which leads to a product which has a relatively poor particle strength.

DE 28 31 561 describes the production of a catalyst support by spray-drying a suspension of pyrolytic silicon dioxide having a concentration of 10 to 33 % and optionally by adding ammonia. The particle size is 40 to 80 μm .

EP 0 341 383 describes a precipitated silicon dioxide with a high "structure", expressed as dibutyl phthalate (DBP) number of 3.0 to 4.0, which corresponds to a high pore volume. In the described process a spray-dryer is used. Given as a possible application is that as matting agents in paints (lacquers).

DE 41 32 230 describes the use of large particles (15 to 100 μm) which were obtained by spray-drying a suspension of milled

silicon dioxide hydrogel. These particles are used as texturing aid for coatings. The used additives have the purpose of producing particles which do not form solid deposits during sedimentation in the liquid paint (lacquer) as a result of standing for a relatively long time. They do not have binding properties. The stability of aggregates decreases with use.

It is the object of the present invention to obtain a matting agent which has an efficiency which is 25 % greater than that of products known from the prior art, without exhibiting a correspondingly greater thickening at the same gloss. Other important properties such as dispersibility, surface roughness and transparency (clarity) of the resulting hardened film are not to be negatively impaired either.

This object is achieved by an aggregated silica gel which has been produced from silica gel particles (particle size: 1 to 20 μ , surface: 200 to 1000 m^2/g , specific pore volume: 0.4 to 2.5 ml/g) and binding agent selected from synthetic or natural layered silicate (phyllosilicate), pyrogenic silicon dioxide and organic polymers soluble in water or dispersible in water.

According to the invention a micronized silica gel having a pore volume of 0.4 to 2.0 ml/g , preferably 1.5 to 1.9 ml/g and in particular 1.6 ml/g , a particle size of 2 to 6 μm , preferably 3 to 4 μm and in particular 4 μm (light diffraction method; Helos Particle Size Analyzer, Sympatec GmbH, Clausthal-Zellerfeld, Germany) and a surface of 200 to 1000 m^2/g , preferably 300 to 450 m^2/g is suspended in water. A binding agent selected from synthetic or natural layered silicate (phyllosilicate), pyrogenic silicon dioxide and organic polymers soluble in water or dispersible in water is dispersed in this suspension using a dissolver for agitation. A surfactant can be added in order to reduce the viscosity and to permit higher solids concentrations. Aggregation takes place in a spray-dryer (Niro, Copenhagen, Type Minor 9). The suspension is atomized in a two-component nozzle

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(suspension throughput: 2 l/h, delivery pressure of the suspension: 500 mm water column, pressure of the compressed (sprayed) air: 2.5 bar above the ambient pressure, quantity of air: 20 m³/h). The water of the formed suspension droplets is evaporated in a hot (inlet temperature: 350°C) counter current air stream. The remaining solid aggregates are removed from the air stream using a cyclone followed by a filter. The coarse particles are then removed from the powder thus formed by screening (classification) in a sifter (Alpine Model MZR). The presence of this coarse fraction (approx. 10 % of the total powder) would give the surface of the paint (lacquer) film a rough undesirable appearance.

Another possibility is to choose the spray conditions such that initially larger aggregates are formed. These are then adjusted to the desired size by milling, followed by classificating (screening). Milling and classificating (screening) can take place at the same time in a jet mill or in separate devices (mill, sifter). The advantage of the desired pore volume increase is retained by this process.

The product according to the invention exhibits a significantly improved matting efficiency compared with the silica gel starting product. This is a result of the additional pore volume (approx. 0.3 to 0.6 ml/g) between the particles of the aggregate (interaggregate volume). The particles themselves have their own internal pore volume.

A certain stability of the aggregate is necessary in order to withstand shear forces during dispersion of the matting agent into the paint. A comparison of the product according to the invention with a standard silicon dioxide matting agent (SYLOID ED5) gives comparable results in this respect (Table 5).

The function of the binding agent is to stabilize the aggregates. Although it is possible to aggregate the micronized silicon

dioxide without binding agent, the stability of this type of aggregates is not sufficient (Table 5).

An optimum stability is achieved with 1 to 25 %, preferably 8 to 12 % of a layered silicate (phyllosilicate) such as Optigel SH from Südchemie. In addition to the preferred synthetic silicate, natural (montmorillonite, hectorite) and synthetic layered silicates (such as laponite), pyrogenic (fumed) silicon dioxide and also organic polymers soluble in water or dispersible in water, such as xanthane, carboxymethyl cellulose (CMC) and polyacrylates are effective.

The thickening power of the matting agent according to the invention is lower than that of the standard SYLOID ED5 when compared at concentrations which are required to achieve the same gloss. This is surprising because in general a higher matting efficiency is associated with a higher thickening power (at the same concentrations), and, when comparing two matting agents at concentrations which lead to the same gloss, an at least equal thickening effect is expected. It is assumed that reasons for the lower viscosity are the spherical shape and the narrow distribution of the particles having a steepness of 0.7 to 1.0 (ED5 has a steepness of 1.2 to 1.45). The steepness of the particle size distribution is defined as

$$St = \frac{d_{v,90} - d_{v,10}}{d_{v,50}}$$

($d_{v,x}$ are defined in DIN 66141).

The other effects of the silicon dioxide in the paint (lacquer) are comparable with those of the standard matting agent.

In order to improve the sedimentation properties of the products according to the invention, it can be treated with a wax selected

from the classes of polyethylene waxes or mineral waxes and modifications thereof.

The products according to the invention can also be used as anti-blocking agents for polymeric films, as beer clarification agents, as thickening agents or abrasives in tooth pastes, in coatings for paper, as catalyst supports and for refining of edible oil.

Examples

Example 1 (Product according to the invention)

500 g SYLOID 244 (surface¹: 410 m²/g, specific pore volume²: 1.58 ml/g, particle size according to light diffraction method (Helos): 3.9 µm) were suspended in 5 l deionized water. 50 g Optigel SH from Südchemie were added and a sodium hydroxide solution was used to adjust the pH value to 9. The suspension was then dispersed for 15 minutes using a 50 mm diameter dissolver blade at 2800 rpm. The suspension was left to stand undisturbed overnight. It was then stirred again at 2800 rpm for 5 minutes. Finally, coarse particles were removed by wet screening through a 100 µm mesh. The suspension was atomized (sprayed) using a two-component nozzle in which the drops were generated using compressed air (2.5 bar above atmospheric). An air stream entering in the spray direction having an inlet temperature of 350°C was used to evaporate the water. The dried material was separated in a cyclone. The median of the volume-related particle size distribution (average volume equivalent particle size diameter) $d_{v,50}$ was adjusted by the pressure of the sprayed air to 10 to 14 µm (light diffraction method; Helos).

¹Nitrogen adsorption (BET) DIN 66131

²Determined with the ASAP 2400 from Micromeritics

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The aggregates were then classified in a dynamic classifier (sifter) Alpin model MZR 100. The particle size was adjusted to a value of 9 to 12 μm by controlling the speed of rotation of the classifier (sifter).

This product was dispersed in a nitrocellulose paint (lacquer) and compared with the standard, SYLOID ED5. The results are given in Table 1. The amount of matting agent required to obtain a gloss level of 30 (angle of incidence 60°) was 24 % lower in the case of the product according to the invention.

The material of Example 1 was also used to matt an alkyd paint (lacquer) (see Table 4). It was compared with the standard, SYLOID ED5, and with TS 100, which is one of the most efficient matting agents on the market and was produced by aggregating pyrolytic silicon dioxide. The matting efficiency of the product according to the invention was 18 % better than that of the standard and 7 % better than that of TS 100. More importantly however, the viscosity of the wet (liquid) paint (lacquer) which contained the product of Example 1 was equal to that of the standard SYLOID ED5, but was significantly lower than the viscosity which arises when using TS 100.

Example 2 (Aggregation without binding agent)

500 g SYLOID 244 (the same product as in Example 1) was treated exactly as in Example 1, without adding a binding agent, and spray dried under the same conditions as in Example 1. The results are given in Table 2. The efficiency of the product was not better than that of the standard SYLOID ED5.

Example 3 (Aggregation with organic binding agent)

3 l of water were heated to 80°C and gently agitated and 12 g xanthene were added in small portions. The hot solution was allowed to cool down after the organic thickening agent had

completely dissolved. 398 g SYLOID 244 (the same product as in Example 1) were then added, and the mixture was treated as in Example 1 and spray dried under the same conditions as in Example 1. The results are given in Table 3. The efficiency was 13 % better than that of the standard SYLOID ED5.

Example 4 (Aggregation with pyrolytic silicon dioxide)

287 g SYLOID 244 (the same product as in Example 1) were suspended in 3 l water. 123 g pyrolytic silicon dioxide (Aerosil 200) were added. The suspension was adjusted to a pH value of 9 by adding sodium hydroxide and spray dried under the same conditions as in Example 1. The results are given in Table 3. the efficiency was 16 % better than that of the standard SYLOID ED5.

Example 5 (improved dispersibility)

10 g of the product produced in Example 1 were filled into a plastic bag measuring 10 cm x 10 cm. A 40 kg weight was placed on it for 30 minutes. A sample of the standard SYLOID ED5 was treated in the same way. The dispersibility of the two samples was compared using a standardized dispersibility test (the product is dispersed for 40 seconds in a nitrocellulose paint (lacquer) using a paint shaker (Red Devil); the number of undispersed agglomerates in the dried paint film is used for characterizing the dispersibility). The film which contained the standard matting agent SYLOID ED5 was full of visible agglomerates, whilst the film which was matted with the product according to the invention was almost free of agglomerates. This is a significant improvement because SYLOID ED5 is regarded as one of the most dispersible products on the market (Tables 1 and 4).

Table 1

Performance of the aggregated matting agent

Parameter	Application test in nitrocellulose paint		
	Unit	Standard SYLOID ED5	Example 1
Particle size [d _{v,50}](1)	μm	8.8	9.2
Matting agent required for 30 gloss units (60° angle of incidence) (2)	% wt./wt.	0.62	0.47
Matting agent required for 40 gloss units (85° angle of incidence) (2)	% wt./wt.	0.72	0.45
Viscosity of the paint with 1 % matting agent (3)	mPa.s	86	83
Dispersibility (4) (low shear)		good	good
Surface roughness Ra at the same gloss (5)	μm	0.4	0.4

- (1) measured according to Helos; focal length 50 mm; median of the volume distribution (DIN 66141)
- (2) DIN 67530
- (3) DIN 53211; beaker opening 4 mm
- (4) Method described in Example 5
- (5) DIN 4768

Table 2

Performance of the aggregated matting agent

Parameter	Application test in nitrocellulose paint		
	Unit	Standard SYLOID ED5	Example 2
Particle size [d _{v,50}] (1)	µm	8.8	8.9
Matting agent required for 30 gloss units (60° angle of incidence) (2)	% wt./wt.	0.71	0.72
Matting agent required for 40 gloss units (85° angle of incidence) (2)	% wt./wt.	0.76	0.84
Viscosity of the paint with 1 % matting agent (3)	mPa.S	361	372

(1) measured according to Helos; focal length 50 mm; median of the volume distribution (DIN 66141)

(2) DIN 67530

(3) DIN 53211; beaker opening 4 mm

Table 3

Performance of the aggregated matting agent

Parameter	Application test in nitrocellulose paint			
	Unit	Standard SYLOID ED5	Example 3	Example 4
Particle size [$d_{v,50}$] (1)	μm	8.8	8.6	9.3
Matting agent required for 30 gloss units (60° angle of incidence) (2)	% wt./wt.	0.61	0.53	0.51
required for 40 gloss units (85° angle of incidence) (2)	% wt./wt.	0.65	0.60	0.47
Viscosity of the paint with 1 % matting agent (3)	mPa.s	357	417	402
Surface roughness Ra at the same gloss (4)	μm	0.65	0.60	0.62

(1) measured according to Helos; focal length 50 mm; median of the volume distribution (DIN 66141)

(2) DIN 67530

(3) DIN 53211; beaker opening 4 mm

(4) DIN 4768

Table 4

Performance of the aggregated matting agent

Parameter	Application test in alkyd paint			
	Unit	Standard SYLOID ED5	Example 1	TS 100
Particle size [$d_{v,50}$] (1)	μm	8.8	9.2	9.1
Matting agent required for 30 gloss units (60° angle of incidence) (2)	% wt./wt.	2.25	1.85	2.00
Viscosity of the paint with 1 % matting agent (3)	mPa.s	111	107	142
Dispersibility (4) (low shear)		good	good	poor
Surface roughness Ra at the same gloss (5)	μm	0.3	0.35	0.3

(1) measured according to Helos; focal length 50 mm; median of the volume distribution (DIN 66141)

(2) DIN 67530

(3) DIN 53211; beaker opening 4 mm

(4) Method described in Example 5

(5) DIN 4768

Table 5

Characterization of the particle stability

Particle size measurement according to different dispersion methods in the original paint (Example 1)

The dispersed sample is diluted prior to measurement using MIBK (methylisobutyl ketone)

Product	Median after dispersion in the dissolution apparatus μm	Median after dispersion in the ball mill	Stability index Ratio of column 2/ column 1
ED5	8.85	8.25	0.93
ED5	8.83	8.32	0.94
Example 1	9.24	8.8	0.95
	8.58	8.13	0.95
Example 2 (no binding agent)	9.21	7.56	0.82
	7.79	6.43	0.83

Patent claims

1. Aggregated silica gel which has been produced from silica gel particles having a particle size of 1 to 20 μm , a surface of 200 to 1000 m^2/g and a specific pore volume of 0.4 to 2.5 ml/g and binding agent selected from synthetic or natural layered silicate (phyllosilicate), pyrolytic (pyrogenic) silicon dioxide and organic polymers soluble in water or dispersible in water.
2. Aggregated silica gel according to claim 1 in which the binding agent is a synthetic or natural phyllosilicate which has been used in a ratio of 1 : 99 to 25 : 75 silicate : silica gel.
3. Aggregated silica gel according to claim 1 in which the binding agent is a pyrolytic silicon dioxide which has been used in a ratio of 5 : 95 to 50 : 50 pyrolytic silicon dioxide : silica gel.
4. Aggregated silica gel according to claim 1 in which the binding agent is an organic polymer soluble in water or dispersible in water which has been used in a ratio of 0.5:99.5 to 15:85 organic polymer:silica gel.
5. Aggregated silica gel according to any of claims 1 to 4 which has been treated with 1 to 20 % of a wax selected from polyethylene waxes or modified polyethylene waxes or mineral waxes or modified mineral waxes or mixtures thereof.
6. Method for the production of an aggregated silica gel as defined in any of claims 1 to 5, wherein aggregation takes place in a spray-dryer using a suspension containing 1 to 25 % solids.

7. Method according to claim 6, wherein the spray-dried solids are then air-classified in order to remove coarse particles.
8. Method according to claim 6, wherein the pH value of the suspension to be spray-dried is adjusted to a value of 8 to 10.5.
9. Method according to claim 6, wherein a wax emulsion is added to the suspension prior to the spray-drying.
10. Method according to claim 6, wherein the wax emulsion is sprayed onto the spray-dried product and is then dried.
11. Use of a product as defined in any of claims 1 to 5 or produced according to any of claims 6 to 10 as matting agent for coatings.
12. Use of a product as defined in any of claims 1 to 5 or produced according to any of claims 6 to 10 as antiblocking agent for polymer films.
13. Use of a product as defined in any of claims 1 to 5 or produced according to one of claims 6 to 10 as beer clarifying agent.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 96/01682

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 C09C1/30 C09D7/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C09C C09D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US,A,4 097 302 (SCM CORPORATION) 27 June 1978 see claim 6	1,5
A	GB,A,1 508 992 (UNILEVER) 26 April 1978 see claims 1,4,6	1,11
P,A	EP,A,0 651 601 (DU PONT DE NEMOURS) 3 May 1995 see claim 1	1,11,12
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☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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2 September 1996

Date of mailing of the international search report

10.09.96

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INTERNATIONAL SEARCH REPORT

International Application No
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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,A	<p>DATABASE WPI Week 9609 Derwent Publications Ltd., London, GB; AN 96-085804 XP002012312 & SU,A,707 076 (AS SIBE CATALYSIS INST. & CATALYSTS CONSTR. TECHN. BUR.) , 27 June 1995 see abstract</p>	1,6
A	<p>EP,A,0 442 325 (GRACE GMBH) 21 August 1991 see page 3, line 43-47; claims 1,2,5,17</p>	1,5,6,11

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

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Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US-A-4097302	27-06-78	NONE	
GB-A-1508992	26-04-78	NONE	
EP-A-651601	03-05-95	DE-A- 4336901 JP-A- 7212012	04-05-95 11-08-95
EP-A-442325	21-08-91	DE-C- 4004468 DE-A- 4032619 AT-T- 119929 AU-B- 638544 AU-B- 7093691 CA-A- 2035675 DE-D- 69108071 DE-T- 69108071 US-A- 5221337	22-08-91 16-04-92 15-04-95 01-07-93 15-08-91 15-08-91 20-04-95 20-07-95 22-06-93